

PCT

WORLD INTELLECTUAL PROPERTY ORGANIZATION
International Bureau



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

<p>(51) International Patent Classification ⁶ : C08F 8/00, 212/14</p>	<p>A1</p>	<p>(11) International Publication Number: WO 96/40798 (43) International Publication Date: 19 December 1996 (19.12.96)</p>
<p>(21) International Application Number: PCT/CA96/00369 (22) International Filing Date: 5 June 1996 (05.06.96) (30) Priority Data: 08/482,948 7 June 1995 (07.06.95) US (60) Parent Application or Grant (63) Related by Continuation US 08/482,948 (CIP) Filed on 7 June 1995 (07.06.95) (71) Applicant (for all designated States except US): BALLARD POWER SYSTEMS INC. [CA/CA]; 9000 Glenlyon Park- way, Burnaby, British Columbia V5J 5J9 (CA). (72) Inventors; and (75) Inventors/Applicants (for US only): STONE, Charles [CA/CA]; 411-237 East 4th Avenue, Vancouver, British Columbia V5T 1G6 (CA). STECK, Alfred, E. [CA/CA]; 3956 Westridge Avenue, West Vancouver, British Colum- bia V7V 3H7 (CA).</p>		<p>(74) Agent: REYNEKE, Russell; Two Bentall Centre, Suite 700, 555 Burrard Street, Vancouver, British Columbia V7X 1M8 (CA). (81) Designated States: AU, CA, JP, US, European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE). Published With international search report.</p>
<p>(54) Title: COPOLYMERIC COMPOSITIONS OF TRIFLUOROSTYRENE, SUBSTITUTED TRIFLUOROSTYRENE AND SUBSTITUTED ETHYLENE, AND ION-EXCHANGE MEMBRANES FORMED THEREFROM</p>		
<p>(57) Abstract</p> <p>Polymeric compositions are derived from copolymers of substituted and unsubstituted α,β-trifluorostyrene, with a variety of substituted ethylene monomers. These compositions are suitable for use as membranes, particularly as ion-exchange membranes, and more particularly as solid polymer electrolytes in electrochemical applications, such as, for example, electrochemical fuel cells.</p>		

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AM	Armenia	GB	United Kingdom	MW	Malawi
AT	Austria	GE	Georgia	MX	Mexico
AU	Australia	GN	Guinea	NE	Niger
BB	Barbados	GR	Greece	NL	Netherlands
BE	Belgium	HU	Hungary	NO	Norway
BF	Burkina Faso	IE	Ireland	NZ	New Zealand
BG	Bulgaria	IT	Italy	PL	Poland
BJ	Benin	JP	Japan	PT	Portugal
BR	Brazil	KE	Kenya	RO	Romania
BY	Belarus	KG	Kyrgyzstan	RU	Russian Federation
CA	Canada	KP	Democratic People's Republic of Korea	SD	Sudan
CF	Central African Republic	KR	Republic of Korea	SE	Sweden
CG	Congo	KZ	Kazakhstan	SG	Singapore
CH	Switzerland	LI	Liechtenstein	SI	Slovenia
CI	Côte d'Ivoire	LK	Sri Lanka	SK	Slovakia
CM	Cameroon	LR	Liberia	SN	Senegal
CN	China	LT	Lithuania	SZ	Swaziland
CS	Czechoslovakia	LU	Luxembourg	TD	Chad
CZ	Czech Republic	LV	Larvia	TG	Togo
DE	Germany	MC	Monaco	TJ	Tajikistan
DK	Denmark	MD	Republic of Moldova	TT	Trinidad and Tobago
EE	Estonia	MG	Madagascar	UA	Ukraine
ES	Spain	ML	Mali	UG	Uganda
FI	Finland	MN	Mongolia	US	United States of America
FR	France	MR	Mauritania	UZ	Uzbekistan
GA	Gabon			VN	Viet Nam

COPOLYMERIC COMPOSITIONS OF TRIFLUOROSTYRENE,
SUBSTITUTED TRIFLUOROSTYRENE
AND SUBSTITUTED ETHYLENE,
AND ION-EXCHANGE MEMBRANES FORMED THEREFROM

Cross-Reference To Related Applications

This application is a continuation-in-part of U.S. Patent Application Serial No. 08/442,206 filed May 16, 1995, which is a continuation of U.S. Patent Application Serial No. 08/124,924 filed September 21, 1993, now U.S. Patent No. 5,422,411 issued June 6, 1995, entitled "Trifluorostyrene And Substituted Trifluorostyrene Copolymeric Compositions And Ion-exchange Membranes Formed Therefrom". The '924 application, incorporated herein by reference in its entirety, describes polymeric compositions derived from copolymers of α,β,β -trifluorostyrene with a variety of substituted α,β,β -trifluorostyrenes. This application is also related to U.S. Patent Application Serial No. 08/480,098 filed June 6, 1995 entitled "Substituted Trifluorostyrene Compositions". The latter application, which is also incorporated herein by reference in its entirety, describes copolymers of α,β,β -trifluorostyrene and substituted α,β,β -trifluorostyrenes including sulfonyl fluoride substituted α,β,β -trifluorostyrene monomers that are conveniently hydrolyzed to produce polymeric compositions with ion-exchange moieties. These compositions are suitable for use as membranes, particularly as ion-exchange membranes.

- 2 -

Field Of The Invention

The present invention relates to substituted ethylene-trifluorostyrene based copolymeric compositions. More particularly, the present invention relates to polymeric compositions derived from copolymers of substituted and unsubstituted α,β,β -trifluorostyrene, with a variety of substituted ethylene monomers. These compositions are particularly suitable for use as solid polymer electrolytes in electrochemical applications, such as, for example, electrochemical fuel cells.

Background Of The Invention

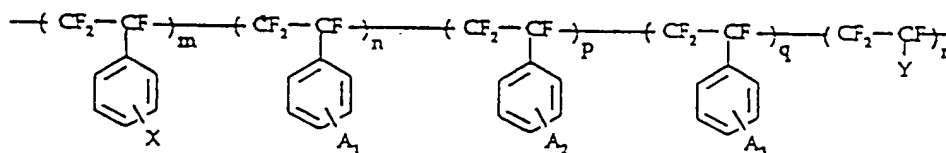
A variety of membranes have been developed over the years for application as solid polymer electrolytes for fuel cells and other electrochemical applications. These polymers have typically been perfluorinated aliphatic compositions, such as those described in U.S. Patent Nos. 3,282,875 and 4,330,654. These compositions are very expensive membranes, and in the case of the '875 patent tend to exhibit poor fuel cell performance characteristic at high current densities. Alternatively, a series of low-cost polyaromatic-based systems have been investigated, such as those described in U.S. Patent Nos. 3,528,858 and 3,226,361. These materials suffer from poor chemical resistance and mechanical properties which tend to limit their use in fuel

- 3 -

cell applications. The investigation of other materials has involved the study of polymers containing the monomer unit α,β,β -trifluorostyrene, for example, those described in U.S. Patent No. 3,341,366 and Japanese Unexamined Patent Publication (Kokai) No. 53-26884. However, these compositions suffered from poor mechanical properties in the case of the '366 patent, and very low polymer yield in the case of the Japanese patent publication.

Summary Of The Invention

Polymeric compositions of the present invention include:

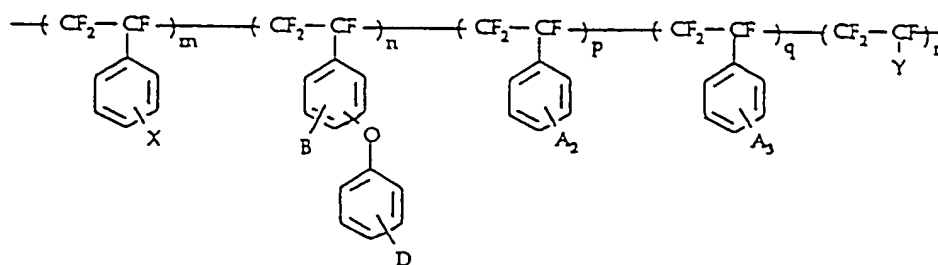


where r is an integer greater than zero, and at least one of m , n , p and q is an integer greater than zero; X is selected from the group consisting of SO_3H , PO_2H_2 , PO_3H_2 , $\text{CH}_2\text{PO}_3\text{H}_2$, COOH , OSO_3H , OPO_2H_2 , OPO_3H_2 , NR_3^+ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls) and CH_2NR_3^+ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls); A_1 , A_2 and A_3 are selected from the group consisting of SO_2F , halogens, alkyls, perfluoroalkyls, O-R (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls), $\text{CF}=\text{CF}_2$, CN , NO_2 and OH ; and Y is selected from the group consisting

- 4 -

of hydrogen, halogens, $C_xH_yF_z$ (where x is an integer greater than zero and $y+z = 2x+1$), $O-R$ (where R is selected from the group consisting of $C_xH_yF_z$ (where x is an integer greater than zero and $y+z = 2x+1$) and aryls), $CF=CF_2$, CN , $COOH$, CO_2R^1 (where R^1 is selected from the group consisting of alkyls, perfluoroalkyls, aryls, and NR^2R^3 where R^2 and R^3 are selected from the group consisting of hydrogen, alkyls and aryls). The group from which A_1 , A_2 and A_3 are selected may further consist of hydrogen provided that Y is not a fluorine, carboxylic acid or ester moiety, or provided that r and at least two of n , p and q are integers greater than zero, or provided that r and m are integers greater than zero.

In an alternative aspect, polymeric compositions of the present invention include:

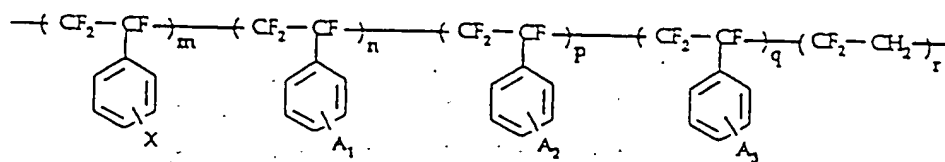


where n and r are integers greater than zero, and m , p and q are zero or an integer greater than zero; X is selected from the group consisting of SO_3H , PO_2H_2 , PO_3H_2 , $CH_2PO_3H_2$, $COOH$, OSO_3H , OPO_2H_2 , OPO_3H_2 , NR_3^+ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls) and $CH_2NR_3^+$ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls); B and D are selected from the group consisting of

- 5 -

hydrogen, SO_2F , SO_3H , PO_2H_2 , PO_3H_2 , $\text{CH}_2\text{PO}_3\text{H}_2$, COOH , OSO_3H , OPO_2H_2 , OPO_3H_2 , NR_3^+ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls) and CH_2NR_3^+ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls); A_2 and A_3 are selected from the group consisting of hydrogen, SO_2F , halogens, alkyls, perfluoroalkyls, O-R (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls), $\text{CF}=\text{CF}_2$, CN, NO_2 , and OH; and Y is selected from the group consisting of hydrogen, halogens, $\text{C}_x\text{H}_y\text{F}_z$ (where x is an integer greater than zero and $y+z = 2x+1$), O-R (where R is selected from the group consisting of $\text{C}_x\text{H}_y\text{F}_z$ (where x is an integer greater than zero and $y+z = 2x+1$) and aryls), $\text{CF}=\text{CF}_2$, CN, COOH , CO_2R^1 (where R^1 is selected from the group consisting of alkyls, perfluoroalkyls, aryls, and NR^2R^3 where R^2 and R^3 are selected from hydrogen, alkyls and aryls).

In a further alternative aspect, polymeric compositions of the present invention include:

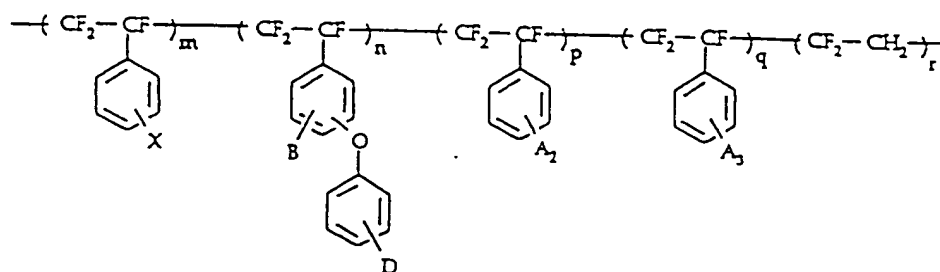


where r is an integer greater than zero, and at least one of m, n, p and q is an integer greater than zero; X is selected from the group consisting of SO_3H , PO_2H_2 , PO_3H_2 , $\text{CH}_2\text{PO}_3\text{H}_2$, COOH , OSO_3H , OPO_2H_2 , OPO_3H_2 , NR_3^+ (where R is selected from the group

- 6 -

consisting of alkyls, perfluoroalkyls and aryls) and CH_2NR_3^+ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls); A_1 , A_2 and A_3 are selected from the group consisting of hydrogen, SO_2F , halogens, alkyls, perfluoroalkyls, O-R (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls), $\text{CF}=\text{CF}_2$, CN, NO_2 , and OH.

In a still further aspect, polymeric compositions of the present invention include:

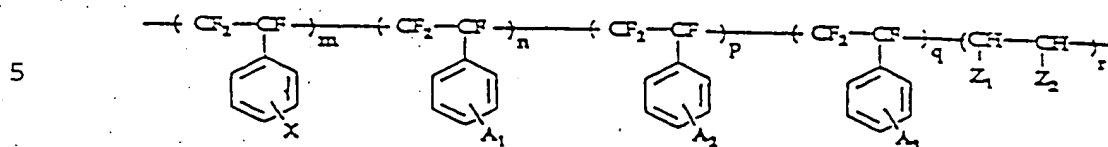


where n and r are integers greater than zero, and m, p and q are zero or an integer greater than zero; X is selected from the group consisting of SO_3H , PO_2H_2 , PO_3H_2 , $\text{CH}_2\text{PO}_3\text{H}_2$, COOH , OSO_3H , OPO_2H_2 , OPO_3H_2 , NR_3^+ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls) and CH_2NR_3^+ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls); B and D are selected from the group consisting of hydrogen, SO_2F , SO_3H , PO_2H_2 , PO_3H_2 , $\text{CH}_2\text{PO}_3\text{H}_2$, COOH , OSO_3H , OPO_2H_2 , OPO_3H_2 , NR_3^+ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls) and CH_2NR_3^+ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls); A_2 and A_3 are selected from the group consisting of hydrogen, SO_2F , halogens, alkyls, perfluoroalkyls, O-R (where R is selected from the

- 7 -

group consisting of alkyls, perfluoroalkyls and aryls), $\text{CF}=\text{CF}_2$, CN , NO_2 , and OH .

In another aspect, polymeric compositions of the present invention include:



where r is an integer greater than zero, and at least one of m , n , p and q is an integer greater than zero; X is selected from the group consisting of SO_3H , PO_2H_2 , PO_3H_2 , $\text{CH}_2\text{PO}_3\text{H}_2$, COOH , OSO_3H , OPO_2H_2 , OPO_3H_2 , NR_3^+ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls) and CH_2NR_3^+ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls); and A_1 , A_2 and A_3 are selected from the group consisting of hydrogen, SO_2F , halogens, alkyls, perfluoroalkyls, O-R (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls), $\text{CF}=\text{CF}_2$, CN , NO_2 and OH ; and Z_1 and Z_2 are selected from the group consisting of hydrogen and fluorine. Thus, in these embodiments the substituted ethylene fragment may be $-\text{CH}_2-\text{CH}_2-$, $-\text{CH}_2-\text{CHF}-$, or $-\text{CHF}-\text{CHF}-$.

The A_1 , A_2 and A_3 substituents may be further elaborated by known techniques such as, for example, hydrolysis of the CN group to form COOH or by reduction with common reducing agents (such as, for example, Raney nickel) to form a primary amine,

- 8 -

thereby transforming the A₁, A₂ and A₃ substituents into ion-exchange moieties.

In any of the embodiments described above, the group from which A₁, A₂ and A₃ are selected can optionally further consist of SO₃H, PO₂H₂, PO₃H₂, CH₂PO₃H₂, COOH, OSO₃H, OPO₂H₂, OPO₃H₂, NR₃⁺ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls) and CH₂NR₃⁺ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls). The resulting polymeric compositions may thus comprise one or more type of ion-exchange moiety, and may also comprise both cation-exchange and anion-exchange moieties.

The substituents on the aromatic rings (A₁, A₂, A₃, X, B and D) may each be located in the ortho, meta or para positions, as indicated in the formulas wherein the chemical bond drawn for these substituents intersects the aromatic ring.

As used herein, the term "aryl" refers to a substituted or unsubstituted aromatic group.

The polymeric compositions of the present invention can also consist essentially of the above chemical units. Thus, the polymers could include amounts of other monomers such as, for example, styrene.

Crosslinking is preferably introduced into the polymeric compositions of the present invention for applications in which it is, for example, desirable to increase dimensional stability, reduce swelling, modify the mechanical properties, or control ion-exchange selectivity.

In accordance with convention in the art, the above chemical formulas for polymeric compositions

- 9 -

containing more than two monomers (where at least three of m, n, p, q and r are greater than zero) are intended to indicate that the monomers are present in the polymeric composition, but are not limited to the particular order in which the monomers are set forth in each general formula. For example, random linear copolymers and/or linear block copolymers formed from the indicated monomers are both contemplated.

The polymeric compositions of the present invention are suitably formed into membranes, and are preferably employed as ion-exchange membranes, most preferably as cation exchange membranes in electrochemical fuel cells.

Brief Description of the Drawing

FIG. 1 is a plot of cell voltage as a function of current density (expressed in amperes per square foot or "ASF") in an electrochemical fuel cell employing, respectively, a Nafion 117 (DuPont's trade designation) cation exchange membrane, a sulfonated α,β,β -trifluorostyrene - tetrafluoroethylene copolymeric membrane (designated "(TFS-TFE)S05"), and a Dow experimental cation exchange membrane (designated "Dow 11").

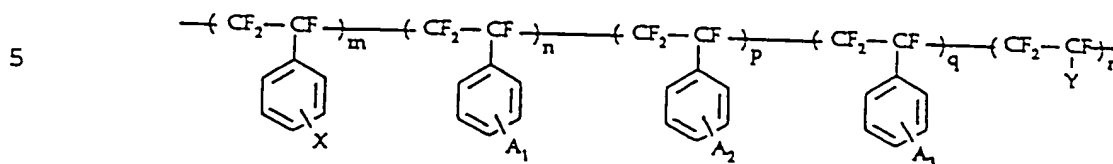
Detailed Description Of The Preferred Embodiments

The present invention relates to polymeric compositions derived from copolymers of substituted and unsubstituted α,β,β -trifluorostyrene, with a variety of substituted ethylene monomers. The substituted ethylenes of the present invention include ethylene itself ($\text{CH}_2=\text{CH}_2$), and tetrafluoroethylene ($\text{CF}_2=\text{CF}_2$), as well as partially

- 10 -

fluorinated ethylenes such as $\text{CH}_2=\text{CHF}$, $\text{CHF}=\text{CHF}$, $\text{CF}_2=\text{CH}_2$, and $\text{CF}_2=\text{CHF}$.

In one aspect, polymeric compositions of the present invention include:

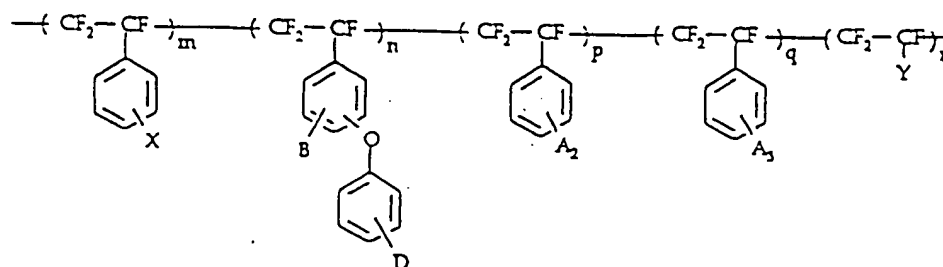


where r is an integer greater than zero, and at least one of m , n , p and q is an integer greater than zero; X is selected from the group consisting of SO_3H , PO_2H_2 , PO_3H_2 , $\text{CH}_2\text{PO}_3\text{H}_2$, COOH , OSO_3H , OPO_2H_2 , OPO_3H_2 , NR_3^+ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls) and CH_2NR_3^+ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls); A_1 , A_2 and A_3 are selected from the group consisting of SO_2F , halogens, alkyls, perfluoroalkyls, O-R (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls), $\text{CF}=\text{CF}_2$, CN , NO_2 and OH ; and Y is selected from the group consisting of hydrogen, halogens, $\text{C}_x\text{H}_y\text{F}_z$ (where x is an integer greater than zero and $y+z = 2x+1$), O-R (where R is selected from the group consisting of $\text{C}_x\text{H}_y\text{F}_z$ (where x is an integer greater than zero and $y+z = 2x+1$) and aryls), $\text{CF}=\text{CF}_2$, CN , COOH , CO_2R^1 (where R^1 is selected from the group consisting of alkyls, perfluoroalkyls, aryls, and NR^2R^3 where R^2 and R^3 are selected from the group consisting of hydrogen, alkyls and aryls). The group from which A_1 , A_2 and A_3 are selected may further consist of hydrogen

- 11 -

provided that Y is not a fluorine, carboxylic acid or ester moiety, or provided that r and at least two of n, p and q are integers greater than zero, or provided that r and m are integers greater than zero.

In an alternative aspect, polymeric compositions of the present invention include:



where n and r are integers greater than zero, and m, p and q are zero or an integer greater than zero; X is selected from the group consisting of SO_3H , PO_2H_2 , PO_3H_2 , $\text{CH}_2\text{PO}_3\text{H}_2$, COOH , OSO_3H , OPO_2H_2 , OPO_3H_2 , NR_3^+ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls) and CH_2NR_3^+ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls); B and D are selected from the group consisting of hydrogen, SO_2F , SO_3H , PO_2H_2 , PO_3H_2 , $\text{CH}_2\text{PO}_3\text{H}_2$, COOH , OSO_3H , OPO_2H_2 , OPO_3H_2 , NR_3^+ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls) and CH_2NR_3^+ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls); A_2 and A_3 are selected from the group consisting of hydrogen, SO_2F , halogens, alkyls, perfluoroalkyls, O-R (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls), $\text{CF}=\text{CF}_2$, CN , NO_2 , and OH ; and Y is selected

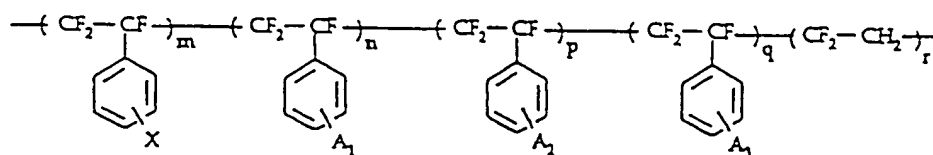
- 12 -

from the group consisting of hydrogen, halogens, $C_xH_yF_z$ (where x is an integer greater than zero and $y+z = 2x+1$), O-R (where R is selected from the group consisting of $C_xH_yF_z$ (where x is an integer greater than zero and $y+z = 2x+1$) and aryls),

5 $CF=CF_2$, CN, COOH, CO_2R^1 (where R^1 is selected from the group consisting of alkyls, C_yF_{2y+1} where y is an integer greater than zero, aryls, and NR^2R^3 where R^2 and R^3 are selected from hydrogen, alkyls and

10 aryls).

In a further alternative aspect, polymeric compositions of the present invention include:



where r is an integer greater than zero, and at

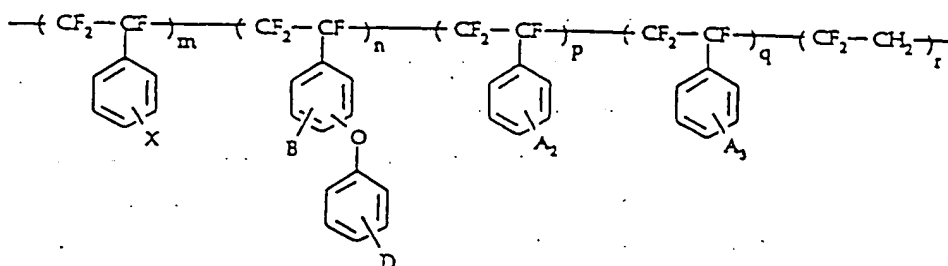
15 least one of m , n , p and q is an integer greater than zero; X is selected from the group consisting of SO_3H , PO_2H_2 , PO_3H_2 , $CH_2PO_3H_2$, COOH, OSO_3H , OPO_2H_2 , OPO_3H_2 , NR_3^+ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls)

20 and $CH_2NR_3^+$ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls); A_1 , A_2 and A_3 are selected from the group consisting of hydrogen, SO_2F , halogens, alkyls, perfluoroalkyls, O-R (where R is selected from the

25 group consisting of alkyls, perfluoroalkyls and aryls), $CF=CF_2$, CN, NO_2 , and OH.

- 13 -

In a still further aspect, polymeric compositions of the present invention include:



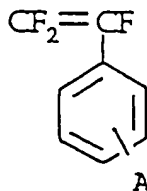
where n and r are integers greater than zero, and m , p and q are zero or an integer greater than zero; X is selected from the group consisting of SO_3H , PO_2H_2 , PO_3H_2 , $\text{CH}_2\text{PO}_3\text{H}_2$, COOH , OSO_3H , OPO_2H_2 , OPO_3H_2 , NR_3^+ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls) and CH_2NR_3^+ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls); B and D are selected from the group consisting of hydrogen, SO_2F , SO_3H , PO_2H_2 , PO_3H_2 , $\text{CH}_2\text{PO}_3\text{H}_2$, COOH , OSO_3H , OPO_2H_2 , OPO_3H_2 , NR_3^+ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls) and CH_2NR_3^+ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls); A_2 and A_3 are selected from the group consisting of hydrogen, SO_2F , halogens, alkyls, perfluoroalkyls, O--R (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls), CF=CF_2 , CN , NO_2 , and OH .

In any of the embodiments described above, the group from which A_1 , A_2 and A_3 are selected can optionally further consist of SO_3H , PO_2H_2 , PO_3H_2 , $\text{CH}_2\text{PO}_3\text{H}_2$, COOH , OSO_3H , OPO_2H_2 , OPO_3H_2 , NR_3^+ (where R is selected from the group consisting of alkyls,

- 14 -

perfluoroalkyls and aryls) and CH_2NR_3^+ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls). The resulting polymeric compositions may thus comprise one or more type of ion-exchange moiety, and may also comprise both cation-exchange and anion-exchange moieties.

Polymeric compositions of the present invention can be produced by polymerization of a substituted ethylene monomer with other monomers selected from a group of substituted α,β,β -trifluorostyrenes having the chemical formula:



In embodiments in which the substituted ethylene monomer is tetrafluoroethylene, A is selected from the group consisting of SO_2F , halogens, alkyls, perfluoroalkyls, O-R (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls), $\text{CF}=\text{CF}_2$, CN, NO_2 and OH.

In embodiments in which the substituted ethylene is 1,1-difluoroethylene, A is selected from the group consisting of SO_2F , halogens, alkyls, perfluoroalkyls, O-R (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls), $\text{CF}=\text{CF}_2$, CN, NO_2 and OH. In an alternative embodiment in which the substituted ethylene is 1,1-difluoroethylene, the group from which A is selected further consists of hydrogen.

- 15 -

In a preferred method, the above monomers are mixed in an aqueous medium containing a free radical initiator and an emulsifying agent, at temperatures in the range of about 35°C - 100°C, and preferably in the range of 45°C - 65°C, for a time period of about 24 to 74 hours under an inert atmosphere. In general, the polymerization procedures and techniques employed in the preparation of polymeric compositions of the present invention are known. A suitable reference for polymerization techniques is Textbook Of Polymer Science, 3rd Edition, by F.W. Billmeyer, Jr., published by John Wiley & Sons.

In general, the monomers used in the preparation of the polymeric compositions of the present invention are commercially available or can be prepared by conventional techniques well known in the art.

Ion-exchange moieties can be introduced into copolymers containing unsubstituted α,β,β -trifluorostyrene units (so-called "base copolymers") via aromatic substitution of at least a portion of those units. For example, base copolymers incorporating pendant unsubstituted phenyl rings can be sulfonated, or in accordance with a further aspect of this invention, may be phosphorylated, carboxylated, quaternary-aminoalkylated or chloromethylated, and further modified to yield $-\text{CH}_2\text{PO}_3\text{H}_2$, $-\text{CH}_2\text{NR}_3^+$ where R is an alkyl, or $-\text{CH}_2\text{NAr}_3^+$ where Ar is a substituted or unsubstituted aromatic moiety, and other substituents, to provide cation-exchange or anion-exchange moieties. Further still, the pendent phenyl moiety may contain a hydroxyl group which can be readily elaborated by

- 16 -

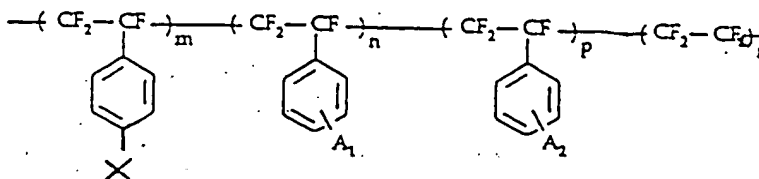
existing methods to generate $-\text{OSO}_3\text{H}$, $-\text{OPO}_3\text{H}_2$, $-\text{OPO}_3\text{H}$ and $-\text{OPO}_3\text{H}_2$ cationic exchange sites on the polymer.

In a typical sulfonation reaction used to produce a cationic exchange membrane, the copolymer is dissolved in an appropriate solvent and then reacted with a sulfonating reagent, such as chlorosulfonic acid or a Lewis acid-base complex of sulfur trioxide. The solvent for such a reaction can be selected from the class consisting of chlorinated aliphatic hydrocarbons, such as dichloroethane, tetrachloroethylene and chloroform. The copolymer solution is rendered completely homogeneous prior to the addition of the solution containing the sulfonating reagent. The reaction is then run within the temperature range from about 10°C up to the boiling point of the solvent, and preferably within the temperature range 18°C - 40°C . To ensure adequate functionalization of the copolymer, the reaction is allowed to continue for a period of about one to about four hours, or longer, dependent on the reaction temperature.

An alternative method of introducing the $-\text{SO}_3\text{H}$ cation exchange moiety is through hydrolysis of $-\text{SO}_2\text{F}$ substituents in the copolymer. In a typical hydrolysis reaction, the sulfonyl fluoride is converted to the free sulfonic acid functionality by hydrolysis in concentrated aqueous alkali metal hydroxide at elevated temperatures. This and other procedures for the hydrolysis of $-\text{SO}_2\text{F}$ to $-\text{SO}_3\text{H}$ are well-known to those skilled in the art.

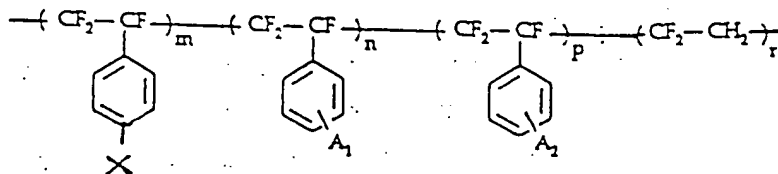
- 17 -

Preferred polymeric compositions of the present invention include:



where m, n and r are integers greater than zero, and p is zero or an integer greater than zero; X is selected from the group consisting of SO₂F and SO₃H; A₁ and A₂ are selected from the group consisting of hydrogen, fluorine, CF₃, and para-phenoxy.

Preferred polymeric compositions of the present invention also include:



where m, n and r are integers greater than zero, and p is zero or an integer greater than zero; X is selected from the group consisting of SO₂F and SO₃H; A₁ and A₂ are selected from the group consisting of hydrogen, fluorine, CF₃, and para-phenoxy.

The substituents on the aromatic rings (A₁, A₂, A₃, X, B and D) in the embodiments described above may be located in the ortho, meta or para positions. In preferred aspects of the described embodiments, the substituents are in the meta or

- 18 -

para positions.

The copolymers thus prepared possess favorable properties, such as thermal stability, chemical resistance and favorable mechanical properties, such as tensile strength, compared to the homopolymeric material formed from α,β,β -trifluorostyrene (TFS) alone.

Crosslinking can be introduced using conventional techniques well-known to those skilled in the art, such as those employed in preparing divinylbenzene crosslinked polystyrene. Crosslinking, for example to enhance the mechanical and physical properties of the membrane material, can be introduced by reaction of appropriate groups, before or preferably after the claimed polymeric compositions are formed into membranes. Monomers with substituents on the pendant phenyl rings or on the ethylene moiety which are suitable for subsequent crosslinking can be introduced into the copolymer in controlled amounts, thereby permitting some control of the degree of crosslinking in the membrane.

The following examples are for purposes of illustration and are not intended to limit the invention. Examples 1 and 2 describe the synthesis of ternary and quaternary copolymers respectively, without ion-exchange moieties, wherein one of the monomers is tetrafluoroethylene. Examples 3 and 4 describe generalized procedures which may be used to prepare copolymers of the present invention. Example 3 describes a suitable method for copolymerization of monomers all of which are either liquid or solids at ambient temperature and pressure. Example 4 describes a suitable method

- 19 -

for copolymerization of monomers, one or more of which is gaseous at ambient temperature and pressure. Examples 5 and 6 describe the sulfonation of two base copolymers to prepare two of the claimed sulfonated copolymers.

Example 7 sets forth the results of tests performed on an ion-exchange copolymer membrane formed from a sulfonated copolymer of the present invention, in an electrochemical fuel cell.

10

Example 1

Emulsion copolymerization of m-trifluoromethyl- α,β,β -trifluorostyrene, α,β,β -trifluorostyrene and tetrafluoroethylene

15

Into a 1 L Parr® reactor under vacuum, is added (by suction) a slurry containing 350 mL of nitrogen-degassed water, dodecylamine hydrochloride (6.8 g, 27 mmol), potassium persulfate (0.52 g 1.8 mmol) and the following monomer mixture:

20

m-trifluoromethyl- α,β,β -trifluorostyrene (11.3 g, 0.05 mol) and α,β,β -trifluorostyrene (47.4 g, 0.30 mol). The reactor is pressurized to approximately 150 psi with tetrafluoroethylene (15 g, 0.15 mol). The resultant emulsion is heated to 50 °C and kept at this temperature for approximately 72 hours.

25

The mixture is then poured into rapidly stirring potassium hydroxide solution (8 g, 0.14 mol in 500 mL water) at 70°C. The polymer which precipitates is filtered, transferred into a Soxhlet thimble and washed by continuous extraction with refluxing methanol to afford an off-white powder; yield 46.0 g (63%), intrinsic viscosity $[\eta] = 0.80$ dL/g as determined in toluene at 30°C. ¹⁹F-NMR analysis

30

- 20 -

performed on a VARIAN XL-300 NMR instrument using CDCl₃ as solvent is used to confirm incorporation of all three monomers.

Example 2

5 Emulsion copolymerization of m-trifluoromethyl-
 α,β,β -trifluorostyrene, p-sulfonyl fluoride- α,β,β -
 trifluorostyrene, α,β,β -trifluorostyrene and
 tetrafluoroethylene

10 Into a 1 L Parr[®] reactor under vacuum, is
 added (by suction) a slurry containing 350 mL of
 nitrogen-degassed water, dodecylamine hydrochloride
 (6.8 g, 27 mmol), potassium persulfate (0.52 g, 1.8
 mol) and the following monomer mixture:
15 m-trifluoromethyl- α,β,β -trifluorostyrene (6.1 g, 54
 mmol), p-sulfonyl fluoride- α,β,β -trifluorostyrene
 (35 g, 0.146 mol) and α,β,β -trifluorostyrene (8.5
 g, 54 mmol). The reactor is then pressurized to 80
 psi with tetrafluoroethylene (8.1 g, 81 mol). The
20 resultant emulsion is then heated to 50 °C and kept
 at this temperature for approximately 72 hours.
 The mixture is then poured into rapidly stirring
 potassium hydroxide solution (8 g, 0.14 mol in 500
 mL water) at 70C°. The polymer which precipitates
25 is filtered, transferred into a Soxhlet thimble and
 washed by continuous extraction with refluxing
 methanol to afford a yellow powder; yield 52.2 g
 (90%), intrinsic viscosity $[\eta] = 1.14$ dL/g as
 determined in toluene at 30°C. ¹⁹F-NMR analysis
30 performed on a VARIAN XL-300 NMR instrument using
 CDCl₃ as solvent is used to confirm incorporation
 of all four monomers.

- 21 -

Example 3

General Emulsion Copolymerization Procedure
(for monomers which are liquid or solid at ambient
temperature and pressure)

5 To a 12 L reaction vessel equipped with a stirrer,
water-cooled condenser, heating mantle and
temperature controller is added 3.2 L of water.
The water is degassed with nitrogen for
approximately one hour and the reaction is kept
10 under a nitrogen atmosphere throughout.
Dodecylamine hydrochloride (58 g, 0.26 mol) is
added and stirred into the water. At this point
the desired monomers (which may be premixed) are
added to the vessel with stirring, to form an
15 emulsion. The temperature of the emulsion is
increased to 50°C and potassium persulfate (4.42 g
15 mmol) is added. The reaction is allowed to
continue for approximately 72 hours. Subsequently,
2 L of water is added to dilute the emulsion,
20 followed by a solution of potassium hydroxide (80
g, 1.43 mol) dissolved in 2 L of water. The
precipitated polymer is then stirred vigorously for
approximately one hour at 75°C. The mixture upon
cooling is filtered, the filter cake being washed
25 several times with fresh water. Having removed the
majority of the filtrate, the cake is then
transferred into a Soxhlet thimble and washed by
continuous extraction with refluxing methanol to
give a random, linear copolymer of the monomers
30 introduced. The resultant product (typically an
off-white powder) is sufficiently pure for further
elaboration.

- 22 -

Example 4

General Emulsion Copolymerization Procedure
(for reactions in which at least one monomer
component is a gas at ambient temperature and
pressure)

The reaction procedure is essentially as described in Example 3 above, with the exception that the presence of a gaseous monomer necessitates that the reaction be performed in a pressurized vessel. Thus, when for example the ethylene-based monomer is gaseous, the reaction is performed in, for example, a Parr® stainless steel reactor. The reactor is equipped with a central stirring shaft, a thermocouple and gas inlet valve, liquid inlet valve and a sampling tube. Depending on the reactivity of the gaseous monomer relative to the other monomer(s), slow addition of the more reactive monomer(s), can be achieved by fixing a pressure-controlled addition funnel to the reactor.

Example 5

Sulfonation of a α,β,β -Trifluorostyrene -
Tetrafluoroethylene Copolymer

Chloroform (200 mL) is added to a 500 mL 3-neck flask fitted with mechanical stirrer, water-cooled condenser and thermocouple attachments. To this is added in portions 7.5 g of a copolymer- (α,β,β -trifluorostyrene - tetrafluoroethylene), which may be prepared according to the method described in Example 4. On obtaining a homogeneous solution, a sulfonating mixture, comprising sulfur trioxide (7.62 g, 95 mmol) and triethyl phosphate

- 23 -

(4.30 g, 24 mmol) in chloroform is rapidly added with vigorous stirring. The resultant mixture is stirred for a further hour at room temperature before quenching the reaction in deionized water. The precipitate is filtered, washed with deionized water, transferred into a Soxhlet thimble and washed by continuous extraction with refluxing chloroform to afford an orange solid powder; yield approximately 11 g, equivalent weight 393 g/mol, water content 184%.

Example 6

Sulfonation of a α,β,β -Trifluorostyrene -
m-Trifluoromethyl- α,β,β -trifluorostyrene -
Tetrafluoroethylene Copolymer

In a procedure essentially identical to that described in Example 5; 7.50 g of copolymer-(α,β,β -trifluorostyrene - m-trifluoromethyl- α,β,β -trifluorostyrene - tetrafluoroethylene), prepared according to the method described in Example 1, dissolved in 200 mL of chloroform is reacted with a sulfonating mixture, comprising sulfur trioxide (7.98 g; 0.10 mole) and triethyl phosphate (4.31 g; 0.024 mole) in chloroform. The resultant mixture is stirred for one hour at room temperature before quenching the reaction in deionized water. The precipitate is filtered, washed with deionized water, transferred into a Soxhlet thimble and washed by continuous extraction with refluxing chloroform to afford an orange powder; yield approximately 10 g, equivalent weight 389 g/mol.

- 24 -

Example 7

An N,N-dimethylformamide solution of a sulfonated α,β,β -trifluorostyrene - tetrafluoroethylene copolymer (equivalent weight 393 g/mol), prepared according to the method described in Example 5, is cast onto a glass substrate using a casting knife. Controlled evaporation of the solvent under a slightly negative pressure results in formation of a membrane (thickness 0.11 mm). The membrane is bonded on opposite major surfaces to two catalyzed carbon paper electrodes at room temperature under 7,500 pounds of pressure. The membrane electrode assembly ("MEA") is tested in the Ballard Mark IV single cell fuel cell (see U.S. Patent Nos. 4,988,583; 5,108,849; 5,170,124; 5,176,966 and 5,200,278; all incorporated herein by reference in their entirety). The results for the sulfonated copolymer membrane tested (designated (TFS-TFE)S05), a DuPont Nafion 117 cation exchange membrane, and a Dow experimental cation exchange membrane (available under the trade designation XUS 13204.10), are shown in Table 1 below.

- 25 -

Table 1

Performance of (TFS-TFE)S05 Copolymer Membrane in a
Ballard Mark IV Fuel Cell

<u>Amps/ft²</u>		<u>Cell Voltage (V)</u>		
		<u>BAM4G</u>	<u>Nafion 117</u>	<u>DOW</u>
5	100	--	0.840	0.846
	200	0.779	0.790	0.818
	300	0.744	0.740	0.783
	400	0.703	0.690	0.748
10	500	0.662	0.650	0.678
	600	0.634	0.580	0.645
	700	0.595	0.510	0.615
	800	0.560	0.430	0.568
	900	0.514	0.330	0.517
15	1000	0.463	0.150	0.440
	1100	0.414	--	0.342

The following operating conditions applied to
the fuel cell in which the membranes were tested:

Temperature = 70°C, reactant inlet pressure 24
20 psi for both air and H₂, reactant
stoichiometries of 2.0 air and 1 to 1.15 H₂.

FIG. 1 is a polarization plot of voltage as a
function of current density in an electrochemical
fuel cell employing, respectively, a DuPont Nafion
25 117 cation exchange membrane, a Dow experimental
cation exchange membrane (available under the trade
designation XUS 13204.10), and sulfonated α,β,β -
trifluorocstyrene - tetrafluoroethylene copolymeric
membrane designated (TFS-TFE)S05 as prepared in

- 26 -

Example 7 above. As shown in FIG. 1, the sulfonated α,β,β -trifluorostyrene - tetrafluoroethylene copolymeric membrane achieved higher cell voltages than the Nafion 117 membrane at all current densities greater than 300 A/ft², and achieved higher cell voltages than the experimental Dow membrane at current densities greater than 900 A/ft².

Copolymers formed from tetrafluoroethylene, α,β,β -trifluorostyrene and various substituted α,β,β -trifluorostyrenes have been produced in yields greater than 60%.

Copolymers of the present invention have the following additional advantages:

1. Flexibility to introduce a wide variety of different ion-exchange functionalities due to the presence of the pendent aromatic groups.
2. The ability to produce a large series of polymeric materials with different equivalent weights starting from the same base copolymer; another flexibility provided by the pendent aromatic substituents.
3. Processibility, in that these copolymers are soluble in a variety of common solvents, for example, N,N-dimethylformamide, dimethyl sulfoxide and N-methylpyrrolidone.
4. The ability to introduce crosslinking, using conventional techniques, such as those employed in preparing divinylbenzene crosslinked polystyrene, to enhance physical and mechanical

- 27 -

properties.

5. Potentially lower cost in view of the low cost of commercially available substituted ethylene monomers.
6. Improved mechanical properties due to increased backbone flexibility and improved dimensional stability through increased overall hydrophobicity resulting from the incorporation of substituted ethylene moieties.

In addition to the utility of the sulfonated copolymeric membranes described herein as ion-exchange membranes for electrochemical fuel cells, the following further utilities are also contemplated:

1. Proton exchange membrane based water electrolysis, which involves a reverse chemical reaction to that employed in hydrogen/oxygen electrochemical fuel cells.
2. Chloralkali electrolysis, typically involving the electrolysis of a brine solution to produce chlorine and sodium hydroxide, with hydrogen as a by-product.
3. Electrode separators in conventional batteries due to the chemical inertness and high electrical conductivity of the sulfonated copolymer membranes.
4. Ion-selective electrodes, particularly those used for the potentiometric determination of a specific ion such as Ca^{2+} , Na^+ , K^+ and like ions. These copolymers could also be employed as the sensor material for humidity sensors, as

- 28 -

- the electrical conductivity of an ion exchange membrane varies with humidity.
5. Ion-exchange material for separations by ion-exchange chromatography. Typical such applications are deionization and desalination of water (for example, the purification of heavy metal contaminated water), ion separations (for example, rare-earth metal ions, trans-uranium elements), and the removal of interfering ionic species.
6. Ion-exchange membranes employed in analytical preconcentration techniques (Donnan Dialysis). This technique is typically employed in analytical chemical processes to concentrate dilute ionic species to be analyzed.
7. Ion-exchange membranes in electrodialysis, in which membranes are employed to separate components of an ionic solution under the driving force of an electrical current. Electrolysis applications include the industrial-scale desalination of brackish water, preparation of boiler feed make-up and chemical process water, de-ashing of sugar solutions, deacidification of citrus juices, separation of amino acids, and the like.
8. Membranes in dialysis applications, in which solutes diffuse from one side of the membrane (the feed side) to the other side according to their concentration gradient. Separation between solutes is

- 29 -

- obtained as a result of differences in diffusion rates across the membrane arising from differences in molecular size. Such applications include
- 5 hemodialysis (artificial kidneys) and the removal of alcohol from beer.
9. Membranes in gas separation (gas permeation) and pervaporation (liquid permeation) techniques.
- 10 10. Bipolar membranes employed in water splitting and subsequently in the recovery of acids and bases from waste water solutions.

While particular elements, embodiments and

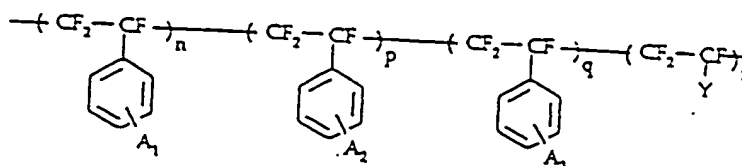
15 applications of the present invention have been shown and described, it will be understood, of course, that the invention is not limited thereto since modifications may be made by those skilled in the art, particularly in light of the foregoing

20 teachings. It is therefore contemplated by the appended claims to cover such modifications as incorporate those features which come within the spirit and scope of the invention.

- 30 -

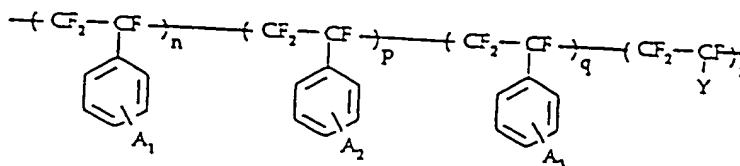
What is claimed is:

1. A polymeric composition comprising:



5 where r is an integer greater than zero, and at least one of n, p and q is an integer greater than zero; A₁, A₂ and A₃ are selected from the group consisting of SO₂F, halogens, alkyls, perfluoroalkyls, O-R (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls), CF=CF₂, CN, NO₂ and OH; and Y is selected from the group consisting of hydrogen, halogens, C_xH_yF_z (where x is an integer greater than zero and y+z = 2x+1), O-R (where R is selected from the group consisting of C_xH_yF_z (where x is an integer greater than zero and y+z = 2x+1) and aryls), CF=CF₂, CN, COOH, CO₂R¹ (where R¹ is selected from the group consisting of alkyls, perfluoroalkyls, aryls, and NR²R³ where R² and R³ are selected from the group consisting of hydrogen, alkyls and aryls).

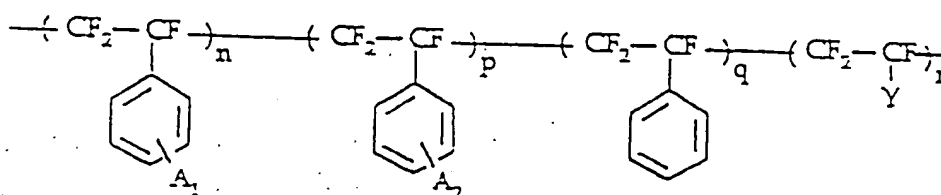
2. A polymeric composition comprising:



- 31 -

where r is an integer greater than zero, and at least one of n , p and q is an integer greater than zero; A_1 , A_2 and A_3 are selected from the group consisting of hydrogen, SO_2F , halogens, alkyls, perfluoroalkyls, O-R (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls), $\text{CF}=\text{CF}_2$, CN , NO_2 and OH ; and Y is selected from the group consisting of hydrogen, $\text{C}_x\text{H}_y\text{F}_z$ (where x is an integer greater than zero and $y+z = 2x+1$), O-R (where R is selected from the group consisting of $\text{C}_x\text{H}_y\text{F}_z$ (where x is an integer greater than zero and $y+z = 2x+1$) and aryls), $\text{CF}=\text{CF}_2$, CN , $\text{CO}_2\text{NR}^2\text{R}^3$ (where R^2 and R^3 are selected from the group consisting of hydrogen, alkyls and aryls).

3. A polymeric composition comprising:

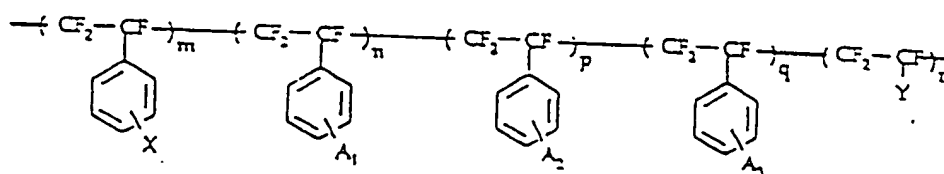


where n , q and r are integers greater than zero, and p is zero or an integer greater than zero; A_1 and A_2 are selected from the group consisting of SO_2F , halogens, alkyls, perfluoroalkyls, O-R (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls), $\text{CF}=\text{CF}_2$, CN , NO_2 and OH ; and Y is selected from the group consisting of hydrogen, halogens, $\text{C}_x\text{H}_y\text{F}_z$ (where x is an integer greater than zero and $y+z = 2x+1$), O-R (where R is selected from the group consisting of $\text{C}_x\text{H}_y\text{F}_z$ (where x is an integer greater than zero and $y+z = 2x+1$) and aryls), $\text{CF}=\text{CF}_2$, CN , COOH , CO_2R^1 (where R^1 is

- 32 -

- 15 selected from the group consisting of alkyls, perfluoroalkyls, aryls and NR^2R^3 where R^2 and R^3 are selected from the group consisting of hydrogen, alkyls and aryls).

4. A polymeric composition comprising:

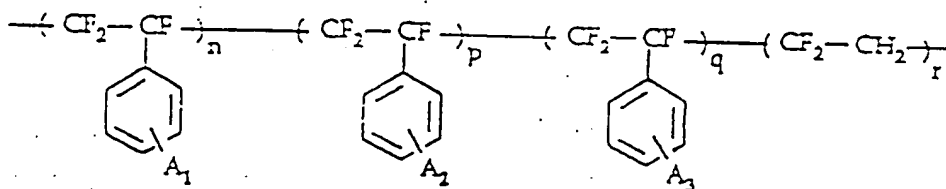


- 5 where m and r are integers greater than zero, and n, p and q are zero or an integer greater than zero; X is selected from the group consisting of SO_3H , PO_2H_2 , PO_3H_2 , $\text{CH}_2\text{PO}_3\text{H}_2$, COOH , OSO_3H , OPO_2H_2 , OPO_3H_2 , NR_3^+ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls) and CH_2NR_3^+ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls);
- 10 A_1 , A_2 and A_3 are selected from the group consisting of hydrogen, SO_2F , halogens, alkyls, perfluoroalkyls, O-R (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls), $\text{CF}=\text{CF}_2$, CN , NO_2 , and OH; and Y is selected
- 15 from the group consisting of hydrogen, halogens, $\text{C}_x\text{H}_y\text{F}_z$ (where x is an integer greater than zero and $y+z = 2x+1$), O-R (where R is selected from the group consisting of $\text{C}_x\text{H}_y\text{F}_z$ (where x is an integer greater than zero and $y+z = 2x+1$) and aryls),
- 20 $\text{CF}=\text{CF}_2$, CN , COOH , CO_2R^1 (where R^1 is selected from the group consisting of alkyls, perfluoroalkyls, aryls, and NR^2R^3 where R^2 and R^3 are selected from the group consisting of hydrogen, alkyls and aryls).

- 33 -

5. The polymeric composition of claim 4 wherein the group from which A_1 , A_2 and A_3 are selected further consists of SO_3H , PO_2H_2 , PO_3H_2 , $\text{CH}_2\text{PO}_3\text{H}_2$, COOH , OSO_3H , OPO_2H_2 , OPO_3H_2 , NR_3^+ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls) and CH_2NR_3^+ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls), and at least one of A_1 , A_2 and A_3 is selected from the group consisting of SO_3H , PO_2H_2 , PO_3H_2 , $\text{CH}_2\text{PO}_3\text{H}_2$, COOH , OSO_3H , OPO_2H_2 , OPO_3H_2 , NR_3^+ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls) and CH_2NR_3^+ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls).

6. A polymeric composition comprising:

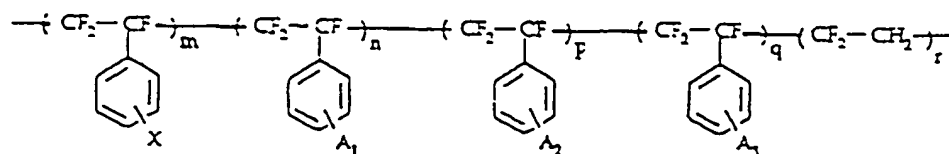


where r is an integer greater than zero, and at least one of n, p and q is an integer greater than zero; A_1 , A_2 and A_3 are selected from the group consisting of SO_2F , halogens, alkyls, perfluoroalkyls, O-R (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls), $\text{CF}=\text{CF}_2$, CN, NO_2 and OH.

7. The polymeric composition of claim 6 wherein the group from which A_1 , A_2 and A_3 are selected further consists of hydrogen.

- 34 -

8. A polymeric composition comprising:



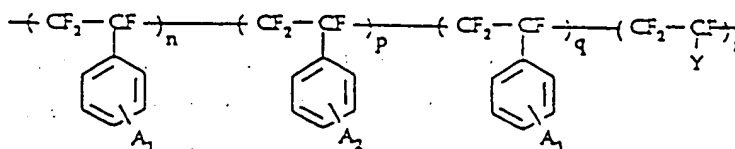
where m and r are integers greater than zero, and
 n, p and q are zero or an integer greater than
 zero; X is selected from the group consisting of
 SO₃H, PO₂H₂, PO₃H₂, CH₂PO₃H₂, COOH, OSO₃H, OPO₂H₂,
 OPO₃H₂, NR₃⁺ (where R is selected from the group
 consisting of alkyls, perfluoroalkyls and aryls)
 and CH₂NR₃⁺ (where R is selected from the group
 consisting of alkyls, perfluoroalkyls and aryls);
 and A₁, A₂ and A₃ are selected from the group
 consisting of hydrogen, SO₂F, halogens, alkyls,
 perfluoroalkyls, O-R (where R is selected from the
 group consisting of alkyls, perfluoroalkyls and
 aryls), CF=CF₂, CN, NO₂, and OH.

9. The polymeric composition of claim 8
 wherein the group from which A₁, A₂ and A₃ are
 selected further consists of SO₃H, PO₂H₂, PO₃H₂,
 CH₂PO₃H₂, COOH, OSO₃H, OPO₂H₂, OPO₃H₂, NR₃⁺ (where R is
 selected from the group consisting of alkyls,
 perfluoroalkyls and aryls) and CH₂NR₃⁺ (where R is
 selected from the group consisting of alkyls,
 perfluoroalkyls and aryls), and at least one of A₁,
 A₂ and A₃ is selected from the group consisting of
 SO₃H, PO₂H₂, PO₃H₂, CH₂PO₃H₂, COOH, OSO₃H, OPO₂H₂,
 OPO₃H₂, NR₃⁺ (where R is selected from the group
 consisting of alkyls, perfluoroalkyls and aryls)

- 35 -

and CH_2NR_3^+ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls).

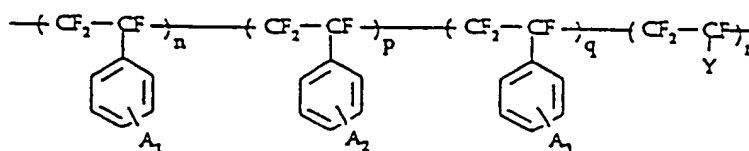
10. A polymeric composition consisting essentially of:



where r is an integer greater than zero, and at least one of n, p and q is an integer greater than zero; A_1 , A_2 and A_3 are selected from the group consisting of SO_2F , halogens, alkyls, perfluoroalkyls, O-R (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls), $\text{CF}=\text{CF}_2$, CN, NO_2 and OH; and Y is selected from the group consisting of hydrogen, halogens, $\text{C}_x\text{H}_y\text{F}_z$ (where x is an integer greater than zero and $y+z = 2x+1$), O-R (where R is selected from the group consisting of $\text{C}_x\text{H}_y\text{F}_z$ (where x is an integer greater than zero and $y+z = 2x+1$) and aryls), $\text{CF}=\text{CF}_2$, CN, COOH, CO_2R^1 (where R^1 is selected from the group consisting of alkyls, perfluoroalkyls, aryls, and NR^2R^3 where R^2 and R^3 are selected from the group consisting of hydrogen, alkyls and aryls).

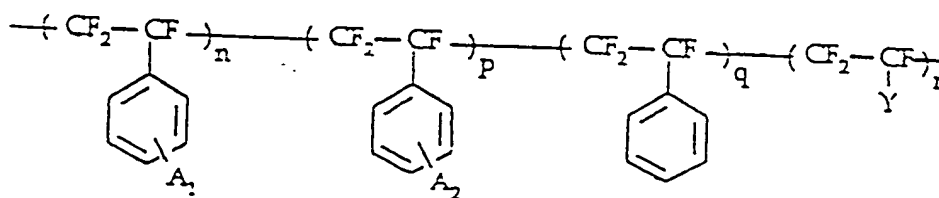
11. A polymeric composition consisting essentially of:

- 36 -



5 where r is an integer greater than zero, and at least one of n, p and q is an integer greater than zero; A₁, A₂ and A₃ are selected from the group consisting of hydrogen, SO₂F, halogens, alkyls, perfluoroalkyls, O-R (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls), CF=CF₂, CN, NO₂ and OH; and Y is selected from the group consisting of hydrogen, C_xH_yF_z (where x is an integer greater than zero and y+z = 2x+1), O-R (where R is selected from the group consisting of C_xH_yF_z (where x is an integer greater than zero and y+z = 2x+1) and aryls), CF=CF₂, CN, CO₂NR²R³ (where R² and R³ are selected from the group consisting of hydrogen, alkyls and aryls).

12. A polymeric composition consisting essentially of:

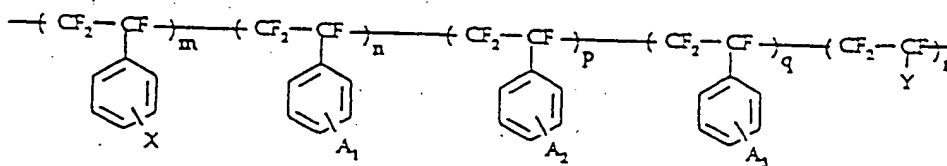


5 where n, q and r are integers greater than zero, and p is zero or an integer greater than zero; A₁ and A₂ are selected from the group consisting of SO₂F, halogens, alkyls, perfluoroalkyls, O-R (where R is selected from the group consisting of alkyls,

- 37 -

- perfluoroalkyls and aryls), $\text{CF}=\text{CF}_2$, CN , NO_2 and OH ;
 10 and Y is selected from the group consisting of hydrogen, halogens, $\text{C}_x\text{H}_y\text{F}_z$ (where x is an integer greater than zero and $y+z = 2x+1$), $\text{O}-\text{R}$ (where R is selected from the group consisting of $\text{C}_x\text{H}_y\text{F}_z$ (where x is an integer greater than zero and $y+z = 2x+1$) and aryls), $\text{CF}=\text{CF}_2$, CN , COOH , CO_2R^1 (where R^1 is selected from the group consisting of alkyls, perfluoroalkyls, aryls and NR^2R^3 where R^2 and R^3 are selected from the group consisting of hydrogen, alkyls and aryls).

13. A polymeric composition consisting essentially of:



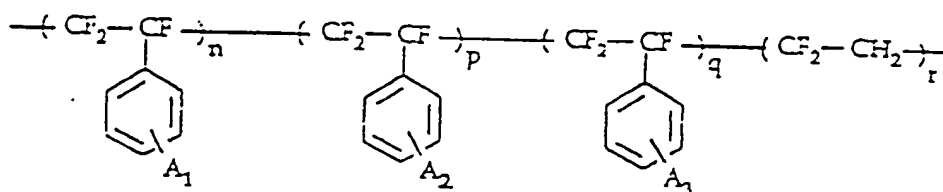
- where m and r are integers greater than zero, and
 5 n, p and q are zero or an integer greater than zero; X is selected from the group consisting of SO_3H , PO_2H_2 , PO_3H_2 , $\text{CH}_2\text{PO}_3\text{H}_2$, COOH , OSO_3H , OPO_2H_2 , OPO_3H_2 , NR_3^+ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls) and CH_2NR_3^+ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls);
 10 A_1 , A_2 and A_3 are selected from the group consisting of hydrogen, SO_2F , halogens, alkyls, perfluoroalkyls, $\text{O}-\text{R}$ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls), $\text{CF}=\text{CF}_2$, CN , NO_2 , and OH ; and Y is selected
 15 from the group consisting of hydrogen, halogens,

- 38 -

20 $C_xH_yF_z$ (where x is an integer greater than zero and $y+z = 2x+1$), O-R (where R is selected from the group consisting of $C_xH_yF_z$ (where x is an integer greater than zero and $y+z = 2x+1$) and aryls), $CF=CF_2$, CN, COOH, CO_2R^1 (where R^1 is selected from the group consisting of alkyls, perfluoroalkyls, aryls, and NR^2R^3 where R^2 and R^3 are selected from the group consisting of hydrogen, alkyls and aryls).

14. The polymeric composition of claim 13 wherein the group from which A_1 , A_2 and A_3 are selected further consists of SO_3H , PO_2H_2 , PO_3H_2 , $CH_2PO_3H_2$, COOH, OSO_3H , OPO_2H_2 , OPO_3H_2 , NR_3^+ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls) and $CH_2NR_3^+$ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls), and at least one of A_1 , A_2 and A_3 is selected from the group consisting of SO_3H , PO_2H_2 , PO_3H_2 , $CH_2PO_3H_2$, COOH, OSO_3H , OPO_2H_2 , OPO_3H_2 , NR_3^+ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls) and $CH_2NR_3^+$ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls).

15. A polymeric composition consisting essentially of:



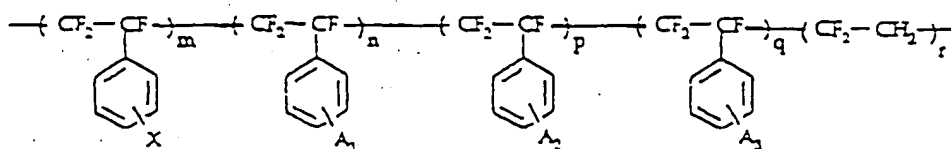
where r is an integer greater than zero, and at

- 39 -

5 least one of n, p and q is an integer greater than zero; A₁, A₂ and A₃ are selected from the group consisting of SO₂F, halogens, alkyls, perfluoroalkyls, O-R (where R is selected from the group consisting of alkyls, perfluoroalkyls and
10 aryls), CF=CF₂, CN, NO₂ and OH.

16. The polymeric composition of claim 15 wherein the group from which A₁, A₂ and A₃ are selected further consists of hydrogen.

17. A polymeric composition consisting essentially of:

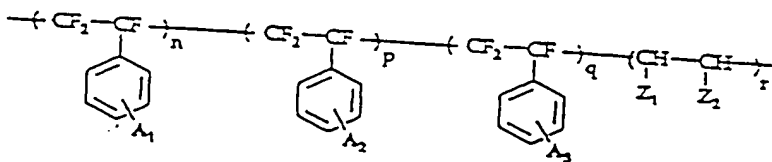


5 where m and r are integers greater than zero, and n, p and q are zero or an integer greater than zero; X is selected from the group consisting of SO₃H, PO₂H₂, PO₃H₂, CH₂PO₃H₂, COOH, OSO₃H, OPO₂H₂, OPO₃H₂, NR₃⁺ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls) and CH₂NR₃⁺ (where R is selected from the group
10 consisting of alkyls, perfluoroalkyls and aryls); and A₁, A₂ and A₃ are selected from the group consisting of hydrogen, SO₂F, halogens, alkyls, perfluoroalkyls, O-R (where R is selected from the group consisting of alkyls, perfluoroalkyls and
15 aryls), CF=CF₂, CN, NO₂, and OH.

- 40 -

18. The polymeric composition of claim 17 wherein the group from which A_1 , A_2 and A_3 are selected further consists of SO_3H , PO_2H_2 , PO_3H_2 , $CH_2PO_3H_2$, $COOH$, OSO_3H , OPO_2H_2 , OPO_3H_2 , NR_3^+ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls) and $CH_2NR_3^+$ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls), and at least one of A_1 , A_2 and A_3 is selected from the group consisting of SO_3H , PO_2H_2 , PO_3H_2 , $CH_2PO_3H_2$, $COOH$, OSO_3H , OPO_2H_2 , OPO_3H_2 , NR_3^+ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls) and $CH_2NR_3^+$ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls).

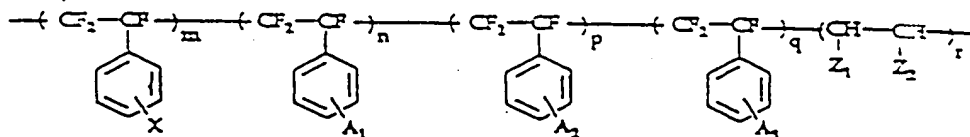
19. A polymeric composition comprising:



where r is an integer greater than zero, and at least one of n , p and q is an integer greater than zero; A_1 , A_2 and A_3 are selected from the group consisting of hydrogen, SO_2F , halogens, alkyls, perfluoroalkyls, $O-R$ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls), $CF=CF_2$, CN , NO_2 and OH ; and Z_1 and Z_2 are selected from the group consisting of hydrogen and fluorine.

20. A polymeric composition comprising:

- 41 -

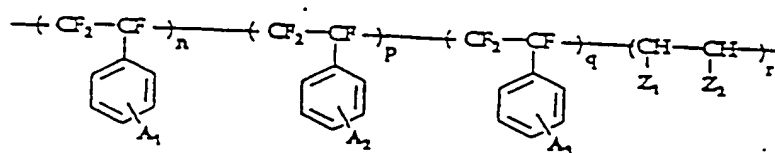


5 where m and r are integers greater than zero, and
 n, p and q are zero or an integer greater than
 zero; X is selected from the group consisting of
 SO₃H, PO₂H₂, PO₃H₂, CH₂PO₃H₂, COOH, OSO₃H, OPO₂H₂,
 OPO₃H₂, NR₃⁺ (where R is selected from the group
 consisting of alkyls, perfluoroalkyls and aryls)
 and CH₂NR₃⁺ (where R is selected from the group
 10 consisting of alkyls, perfluoroalkyls and aryls);
 and A₁, A₂ and A₃ are selected from the group
 consisting of hydrogen, SO₂F, halogens, alkyls,
 perfluoroalkyls, O-R (where R is selected from the
 group consisting of alkyls, perfluoroalkyls and
 aryls), CF=CF₂, CN, NO₂ and OH; and Z₁ and Z₂ are
 15 selected from the group consisting of hydrogen and
 fluorine.

21. The polymeric composition of claim 20
 wherein the group from which A₁, A₂ and A₃ are
 selected further consists of SO₃H, PO₂H₂, PO₃H₂,
 CH₂PO₃H₂, COOH, OSO₃H, OPO₂H₂, OPO₃H₂, NR₃⁺ (where R is
 5 selected from the group consisting of alkyls,
 perfluoroalkyls and aryls) and CH₂NR₃⁺ (where R is
 selected from the group consisting of alkyls,
 perfluoroalkyls and aryls), and at least one of A₁,
 A₂ and A₃ is selected from the group consisting of
 10 SO₃H, PO₂H₂, PO₃H₂, CH₂PO₃H₂, COOH, OSO₃H, OPO₂H₂,
 OPO₃H₂, NR₃⁺ (where R is selected from the group

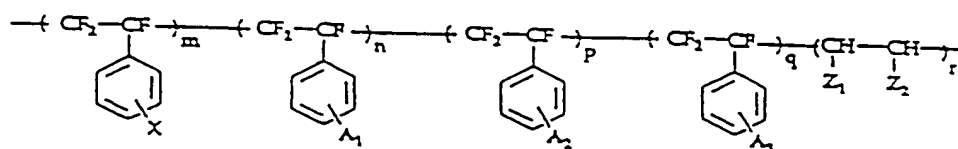
consisting of alkyls, perfluoroalkyls and aryls) and CH_2NR_3^+ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls).

22. A polymeric composition consisting essentially of:



where r is an integer greater than zero, and at least one of n , p and q is an integer greater than zero; A_1 , A_2 and A_3 are selected from the group consisting of hydrogen, SO_2F , halogens, alkyls, perfluoroalkyls, $O-R$ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls), $CF=CF_2$, CN , NO_2 and OH ; and Z_1 and Z_2 are selected from the group consisting of hydrogen and fluorine.

23. A polymeric composition consisting essentially of:



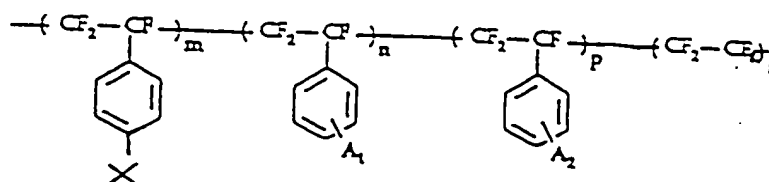
- 43 -

where m and r are integers greater than zero, and n, p and q are zero or an integer greater than zero; X is selected from the group consisting of SO₃H, PO₂H₂, PO₃H₂, CH₂PO₃H₂, COOH, OSO₃H, OPO₂H₂, OPO₃H₂, NR₃⁺ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls) and CH₂NR₃⁺ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls); and A₁, A₂ and A₃ are selected from the group consisting of hydrogen, SO₂F, halogens, alkyls, perfluoroalkyls, O-R (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls), CF=CF₂, CN, NO₂ and OH; and Z₁ and Z₂ are selected from the group consisting of hydrogen and fluorine.

24. The polymeric composition of claim 23 wherein the group from which A₁, A₂ and A₃ are selected further consists of SO₃H, PO₂H₂, PO₃H₂, CH₂PO₃H₂, COOH, OSO₃H, OPO₂H₂, OPO₃H₂, NR₃⁺ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls) and CH₂NR₃⁺ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls), and at least one of A₁, A₂ and A₃ is selected from the group consisting of SO₃H, PO₂H₂, PO₃H₂, CH₂PO₃H₂, COOH, OSO₃H, OPO₂H₂, OPO₃H₂, NR₃⁺ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls) and CH₂NR₃⁺ (where R is selected from the group consisting of alkyls, perfluoroalkyls and aryls).

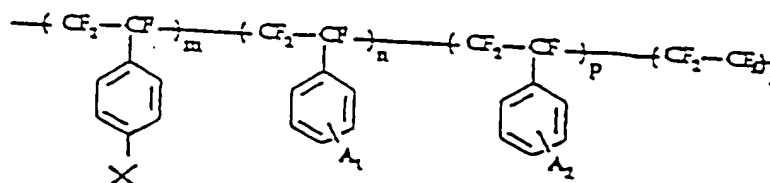
25. A polymeric composition comprising:

- 44 -



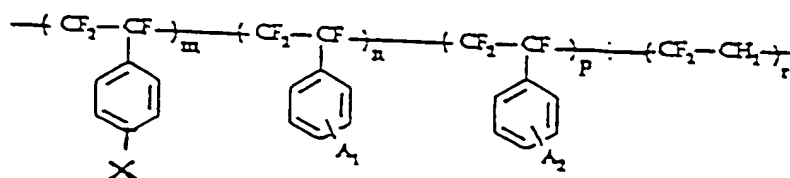
5 where m, n and r are integers greater than zero, and p is zero or an integer greater than zero: X is selected from the group consisting of SO₂F and SO₃H; A₁ and A₂ are selected from the group consisting of hydrogen, fluorine, CF₃, and para-phenoxy.

26. A polymeric composition consisting essentially of:



5 where m, n and r are integers greater than zero, and p is zero or an integer greater than zero: X is selected from the group consisting of SO₂F and SO₃H; A₁ and A₂ are selected from the group consisting of hydrogen, fluorine, CF₃, and para-phenoxy.

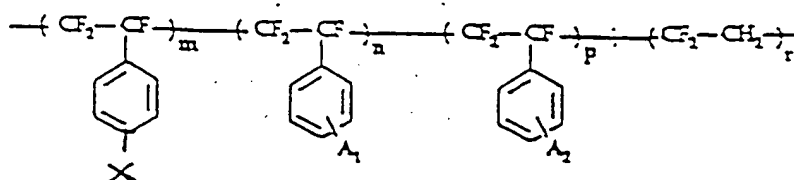
27. A polymeric composition comprising:



- 45 -

where m, n and r are integers greater than zero, and p is zero or an integer greater than zero: X is selected from the group consisting of SO₂F and SO₃H; A₁ and A₂ are selected from the group consisting of hydrogen, fluorine, CF₃, and para-phenoxy.

28. A polymeric composition consisting essentially of:



where m, n and r are integers greater than zero, and p is zero or an integer greater than zero: X is selected from the group consisting of SO₂F and SO₃H; A₁ and A₂ are selected from the group consisting of hydrogen, fluorine, CF₃, and para-phenoxy.

29. The polymeric composition of any of claims 1-28 wherein the composition is at least partially crosslinked.

30. A polymeric membrane comprising the polymeric composition of any of claims 1-29.

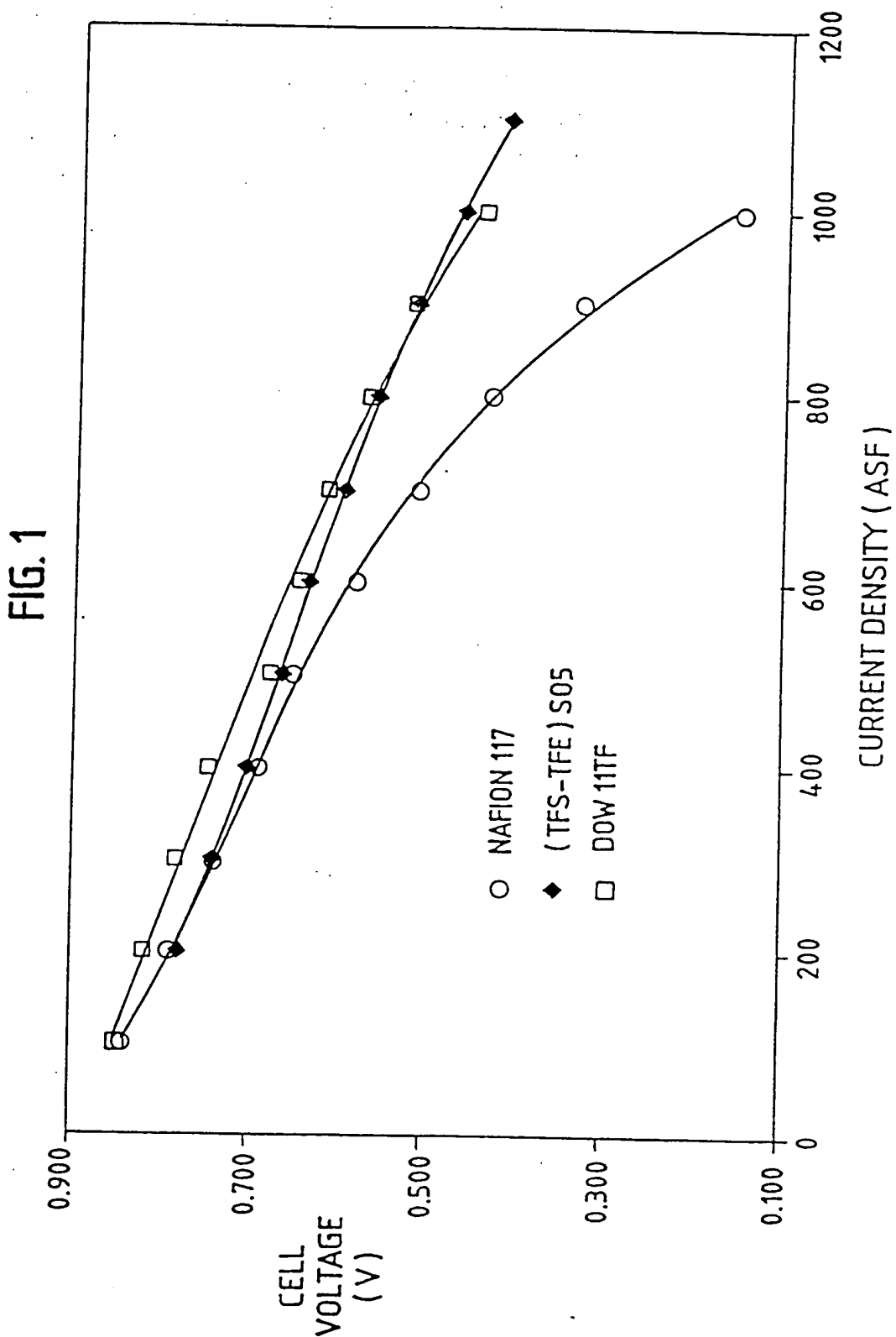
31. A polymeric membrane consisting essentially of the polymeric composition of any of claims 1-29.

- 46 -

32. An electrochemical fuel cell comprising the polymeric membrane of claim 30 wherein said polymeric membrane is an ion exchange membrane.

33. An electrochemical fuel cell comprising the polymeric membrane of claim 31 wherein said polymeric membrane is an ion exchange membrane.

1/1



This Page Blank (uspto)

INTERNATIONAL SEARCH REPORT

International Application No.
PCT/CH 96/00369

A. CLASSIFICATION OF SUBJECT MATTER IPC 6 C08F8/00 C08F212/14		
According to International Patent Classification (IPC) or to both national classification and IPC		
B. FIELDS SEARCHED		
Minimum documentation searched (classification system followed by classification symbols) IPC 6 C08F		
Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched		
Electronic data base consulted during the international search (name of data base and, where practical, search terms used)		
C. DOCUMENTS CONSIDERED TO BE RELEVANT		
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO,A,95 08581 (BALLARD POWER SYSTEMS INC.) 30 March 1995 cited in the application see page 4, line 10 - page 5, line 19 see page 18, line 16 - page 20, line 25; claims 1-18	1-3, 10-12, 29-33
A	--- FR,A,2 096 180 (DUNLOP HOLDINGS LTD.) 11 February 1972 see claims 1-26	1
A	--- US,A,4 126 588 (HIROSHI UKIHASHI) 21 November 1978 see claims 1-13 ---	1
-/--		
<input checked="" type="checkbox"/> Further documents are listed in the continuation of box C. <input checked="" type="checkbox"/> Patent family members are listed in annex.		
* Special categories of cited documents :		
<div style="display: flex; justify-content: space-between;"> <div style="width: 45%;"> <p>"A" document defining the general state of the art which is not considered to be of particular relevance</p> <p>"E" earlier document but published on or after the international filing date</p> <p>"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)</p> <p>"O" document referring to an oral disclosure, use, exhibition or other means</p> <p>"P" document published prior to the international filing date but later than the priority date claimed</p> </div> <div style="width: 45%;"> <p>"T" later document published after the international filing date or priority date and not in conflict with the application but cited to undermine the principle or theory underlying the invention</p> <p>"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone</p> <p>"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.</p> <p>"&" document member of the same patent family</p> </div> </div>		
Date of the actual completion of the international search <div style="text-align: center; font-weight: bold;">18 September 1996</div>	Date of mailing of the international search report <div style="text-align: center; font-weight: bold;">- 3. 10. 96</div>	
Name and mailing address of the ISA European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+ 31-70) 340-2040, Tx. 31 651 epo nl, Fax (+ 31-70) 340-3016	Authorized officer <div style="text-align: center; font-weight: bold;">Permentier, W</div>	

This Page Blank (uspto)

INTERNATIONAL SEARCH REPORT

International Application No.

PCT/CA 96/00369

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	<p>US,A,3 442 825 (R. B. HODGDON) 6 May 1969 cited in the application see column 3, line 62 - column 4, line 22 see column 4, line 64 - column 5, line 15; claims 1-6</p> <p style="text-align: center;">-----</p>	1,29-33

This Page Blank (uspto)

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/CA 96/00369

PC/CA 30/00309

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO-A-9508581	30-03-95	US-A- 5422411	06-06-95
		AU-A- 7649294	10-04-95
		CA-A- 2171298	30-03-95
		EP-A- 0720628	10-07-96
		US-A- 5498639	12-03-96

FR-A-2096180	11-02-72	GB-A- 1352138	08-05-74

US-A-4126588	21-11-78	JP-C- 1226180	31-08-84
		JP-A- 52116790	30-09-77
		JP-B- 58000895	08-01-83
		JP-C- 1078230	25-12-81
		JP-A- 52084191	13-07-77
		JP-B- 53044429	29-11-78
		DE-A- 2659581	14-07-77
		FR-A- 2337162	29-07-77
		GB-A- 1531068	01-11-78

US-A-3442825	06-05-69	BE-A- 668486	16-12-65
		CH-A- 463462	
		DE-A- 1570681	12-03-70
		FR-A- 1452148	08-12-66
		GB-A- 1096879	
		NL-A- 6510859	21-02-66
		SE-B- 337921	23-08-71
		US-A- 3341366	12-09-67

Page Blank (uspto)